

# Mitogen-Activated Protein Kinase Activation by Oxidative and Bacterial Stress in an Amphibian Cell Culture Model

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The decline of many amphibian species could be caused by their susceptibility to environmental pollutants that cause cellular stress and cell death. A variety of intracellular signal transduction pathways are activated by environmental stress factors, which result in cell death. Mitogen-activated protein kinases are intracellular signaling molecules that include the extracellular signal-regulated kinases (ERK-1 and ERK-2). We used cultured *Xenopus* tadpole cells (XTC-2 cells) to investigate the activation of ERK by oxidative or bacterial stress, two environmental factors that could contribute to pollution in aquatic systems. We exposed XTC-2 cell monolayers to hydrogen peroxide or bacterial lipopolysaccharide and measured ERK activation by Western blotting using antibodies raised against phosphorylated ERK-1 and ERK-2. Only ERK-2 was detected in XTC-2 cells. Both hydrogen peroxide and lipopolysaccharide caused ERK-2 phosphorylation in a time- and concentration-dependent manner. Hydrogen peroxide caused a 20- to 30-fold increase in ERK-2 activation that peaked 30 min after treatment, and lipopolysaccharide induced a 5- to 10-fold increase in ERK-2 activation that peaked 60 min after treatment. PD98059, an inhibitor of the ERK pathway, reduced the cytotoxic response of XTC-2 cells to hydrogen peroxide or lipopolysaccharide. These data suggest that ERK-2 is an intracellular target of oxidative and bacterial stress in amphibians that mediates, at least in part, the cytotoxic response to hydrogen peroxide or lipopolysaccharide. Moreover, the *Xenopus* (XTC-2) cell culture system could serve as a useful model to identify agents that might threaten amphibian populations and human health. **Key words:** amphibian decline, endotoxins, environmental stress, mitogen-activated protein kinase, reactive oxygen species, *Xenopus*. *Environ Health Perspect* 110:641–645 (2002). [Online 15 May 2002]

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Amphibians are particularly sensitive to environmental stress due to their unique life-history strategies that lead to their exposure to natural and anthropogenic perturbations in both aquatic and terrestrial ecosystems. Because amphibians have thin, permeable skin and their eggs lack shells, they are susceptible to environmental exposure during development and as adults. Therefore, they may serve as important sentinel species to changes in environmental conditions. Since the mid-1990s, unusually large numbers of frogs with malformations have been found in North America (1,2). Abnormalities have been reported in 36 species of amphibians from 42 U.S. states. There is also concern that the environmental factors causing these anuran deformities may affect human health. The reason for the amphibian malformations and population decline remains unknown, although it has been suggested that parasitic infections, UV radiation, and chemical pollutants could play a role (3). If endoparasitic infestations are involved, it is likely that environmental stress (i.e., UV radiation, pollution) could increase susceptibility to these infections (4). It is also reasonable to assume that a number of critical environmental interactions could contribute to the detrimental effects on amphibian populations (5,6).

There has also been a rapid decline and range reduction of certain amphibian species worldwide over the past two decades. However, some species within the same regions do not appear to be affected, whereas others are driven to extinction (7). Although the explanation for the degradation of amphibian populations is currently disputed, Pechmann et al. (8) caution that it may be difficult to distinguish natural population fluctuations from human-induced declines without long-term surveys at numerous sites to isolate the various confounding components. However, anthropogenic factors such as water pollution and atmospheric conditions such as ozone depletion and/or UV radiation could play important roles (4,6). Continuing population reductions and morphologic defects in anurans has generated international interest. A primary consideration in the correlation of this problem to human health is our dependence on water quality. Therefore, frogs are an important sentinel species that may indicate environmental deterioration. Although there has been much investigation into the causal factors contributing to malformations and declining amphibian numbers, *in vitro* cell culture models to address which environmental pollutants cause cellular stress in amphibians are lacking.

A major indicator of cellular stress is the activation of a class of intracellular proteins called mitogen-activated protein (MAP) kinases, which are a family of serine/threonine kinases that include extracellular signal-regulated kinases (ERKs), p38 MAP kinases, and *c-jun* N-terminal kinases (JNKs) (9). MAP kinases normally regulate a variety of cellular activities in response to endogenous cytokines and growth factors, leading to biologic outcomes such as cell growth and differentiation (10). Several environmental agents activate MAP kinases to cause cellular stress and cytotoxic responses, including bacterial products (such as lipopolysaccharide; LPS), pesticides, certain transition metals, and UV radiation (11–13). Transition metals and UV radiation cause MAP kinase activation via the generation of reactive oxygen species (14,15). In particular, UV radiation has been implicated as a major factor in amphibian mortality and deformities (16–19).

In this study, we investigated MAP kinase (ERK) activation in *Xenopus* tadpole cells (XTC-2). The XTC-2 cells were treated with a reactive oxygen species (hydrogen peroxide; H<sub>2</sub>O<sub>2</sub>) or a bacterial wall product, LPS. Both of these factors are associated with environmental stress in polluted aquatic ecosystems (20,21). H<sub>2</sub>O<sub>2</sub> and LPS both caused phosphorylation of ERK-2 in XTC-2 cells in a time- and dose-dependent manner. These data suggest that the ERK-2 pathway is a sensitive biomarker to detect environmental stress and that the ERK pathway mediates cell death in response to H<sub>2</sub>O<sub>2</sub> and LPS.

## Materials and Methods

**Materials.** Goat polyclonal anti-ERK-1/2 (catalog no. SC-154-G) that recognizes total cellular ERK (i.e., both phosphorylated and unphosphorylated forms) was purchased from Santa Cruz Biotechnology (Santa Cruz, CA). This antibody was raised against an epitope mapping the carboxy terminus of

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ERK-2-encoded MAP kinase p42 of rat origin. The antibody is reactive with ERK-2 and to a lesser extent ERK-1 and possesses broad interspecies reactivity (murine, human, chicken, and frog). To verify that we were detecting only ERK-2 and not ERK-1 in XTC-2 cells, we used two other antibodies raised against total ERK-1 and -2 from Transduction Laboratories (Lexington, KY) or Upstate Biotechnologies (Lake Placid, NY). An anti-phospho-ERK-1/2 monoclonal antibody that is specific for the phosphorylated forms of ERK-1/2 (catalog no. 9106) was purchased from Cell Signaling Technologies/New England Biolabs (Beverly, MA). This antibody had previously been reported by the company to cross-react with ERK-1/2 from human, rat, mouse, and hamster (no information on *Xenopus* was available). Anti-mouse IgG-HRP was purchased from Transduction Laboratories, and anti-goat IgG-HRP was purchased from DakoPatts (Carpenteria, CA). *E. coli* LPS (no. L8274) and  $H_2O_2$  were purchased from Sigma Chemical Company (St. Louis, MO). The MEK/ERK inhibitor, PD98059, was purchased from Calbiochem (La Jolla, CA).

**Cell culture.** *Xenopus* tadpole cells (XTC-2) were kindly provided by P. Blackshear at NIEHS and were originally obtained from D. J. Shapiro at the University of Illinois-Urbana (22). Aliquots of XTC-2 thawed from liquid nitrogen storage were seeded in culture flasks containing Leibovitz's L-15 medium supplemented with 10% fetal bovine serum (Sigma). Cultures of XTC-2 were grown in 175 cm<sup>2</sup> sterile, plastic culture flasks at room temperature with humidified air. At confluence, the cells were liberated

from the culture surface with a 1:10 dilution of trypsin (Gibco, Grand Island, NY) in 1 mM EDTA/phosphate-buffered saline, pH 7.4. Cells were then plated in 75-cm<sup>2</sup> plastic culture dishes, grown to confluence, and growth arrested in serum-free Leibovitz's L-15 medium for 24 hr before treatment with  $H_2O_2$  or LPS.

**Western blotting.** After treating XTC-2 cells with  $H_2O_2$  or LPS, the cells were placed on ice and washed twice with ice-cold phosphate-buffered saline (PBS), and after all the residual PBS was carefully removed, 100  $\mu$ L of lysis buffer was added to the cell monolayer. The lysis buffer consisted of 50 mM HEPES, 150 mM NaCl, 1% Triton X-100, 1 mM PMSF, and 20  $\mu$ g/mL aprotinin, leupeptin, and pepstatin. The cellular protein was then removed from the culture surface with a cell scraper and lysates were clarified by centrifugation at 13,000 rpm for 10 min and frozen at  $-80^\circ\text{C}$ . We loaded 20  $\mu$ g of protein from each sample on an 8–16% SDS-PAGE gel. Separated proteins were then transferred to a nitrocellulose membrane, and the membrane was blocked for 2 hr at room temperature with 5% non-fat milk in Tris-buffered saline-Tween buffer (20 mM Tris, 500 mM NaCl, 0.01% Tween 20). The membrane was then incubated with the anti-phospho-ERK-1/2 antibody (1:1,000 dilution) at  $4^\circ\text{C}$  overnight, followed by incubation for 2 hr with a 1:2,000 dilution of horseradish peroxidase-conjugated anti-mouse IgG. The immunoblot was visualized through enhanced chemiluminescence (Amersham, Arlington Heights, IL). The same blot was subsequently stripped at  $50^\circ\text{C}$  for 30 min in a buffer containing 62.5 mM Tris (pH 6.7), 2% SDS, and 100 mM 2-mercaptoethanol. The stripped blots were incubated with polyclonal anti-ERK-1/2 (1:1,000 dilution) at  $4^\circ\text{C}$  overnight, followed by incubation for 2 hr with a 1:2,000 dilution

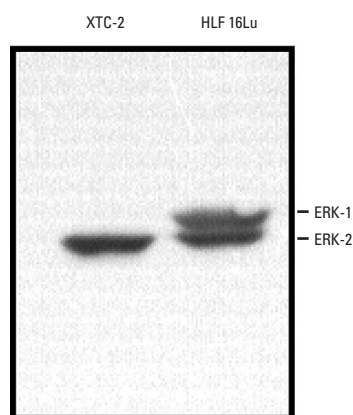
of horseradish peroxidase-conjugated anti-goat IgG.

**Cytotoxicity assay.** We measured cytotoxicity by the tetrazolium salt (MTT) cell viability assay according to the manufacturer's instructions (Boehringer Mannheim, Indianapolis, IN). Absorbance was measured at 570 nm with a 630 nm reference filter. The negative control was the absorbance of cells with medium alone. The positive control (100% cytotoxicity) was determined by treating cells with distilled water for 24 hr.

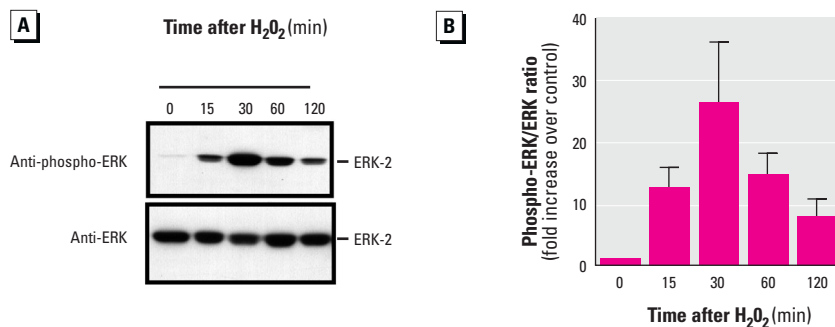
**Densitometric evaluation of ERK activation and statistical analysis.** Autoradiographic images from the Western blots were digitally scanned and densitometric measurements for phospho-ERK and total ERK signals were performed using the NIH Image program. The phospho-ERK signal was normalized against the corresponding total ERK signal from the same Western blot to obtain the phospho-ERK/ERK ratio. In all experiments, Western blotting was performed first for phospho-ERK (Cell Signaling, catalog no. 9106), then the blots were stripped and reblotted for total ERK using an antibody that recognizes the total ERK (Santa Cruz, catalog no. SC-154-G). The data were expressed as fold-increase over control (i.e., cells treated with medium alone). We repeated each experiment at least three times and determined significant differences by one-way analysis of variance (ANOVA). For cytotoxicity data, we analyzed comparisons of control versus treatment by paired *t*-tests.

## Results

**XTC-2 cells contain ERK-2.** We investigated MAP kinase activation in confluent, quiescent cultures of *Xenopus* (XTC-2) cells as an *in vitro* model of oxidant- and bacterial-induced cellular stress. We first determined whether XTC-2 cells possessed either the p42 (ERK-2) or p44 (ERK-1) isozymes of MAP kinase.



**Figure 1.** Expression of the MAP kinase ERK-2 in amphibian cells (XTC-2) and in human lung fibroblasts (HLF). Cell lysates from confluent, quiescent cultures of XTC-2 cells and HLF cells were assayed by Western blotting using an anti-ERK antibody that detects unphosphorylated ERK-1 and ERK-2. XTC-2 cells contained only ERK-2 whereas both ERK-1 and ERK-2 were present in HLF cells.



**Figure 2.** Western blot analysis showing phosphorylation of ERK-2 by  $H_2O_2$  in a time-dependent manner. Confluent, quiescent XTC-2 cells were treated with  $H_2O_2$  (10  $\mu$ M) for 15, 30, 60, and 120 min. (A) Representative Western blot experiment showing that maximum phosphorylation of ERK-2 was observed at 30 min after treatment with  $H_2O_2$ . (B) Relative levels of ERK-2 activation determined by densitometric scanning of the phospho-ERK bands and normalized to the ERK signal. Data are the means  $\pm$  SEM of three separate experiments and are presented as the phospho-ERK-2/ERK-2 ratio. One-way ANOVA yielded a calculated *F* value of 3.12 ( $p = 0.049$ ).

We detected only ERK-2 (p42 MAP kinase) in XTC-2 cells, whereas both ERK-1 and ERK-2 were readily detectable in human lung fibroblasts (Figure 1). Similar results were obtained using two other polyclonal antibodies raised against total ERK-1 and ERK-2 (data not shown). Therefore, although we cannot rule out that XTC-2 cells might possess ERK-1, we clearly demonstrated that our assay detects only ERK-2.

**Rapid activation of ERK-2 by oxidative stress.** We next evaluated whether oxidative stress would activate ERK-2 in XTC-2 cells, as has been reported in mammalian cell culture systems (23). ERK-2 phosphorylation peaked 30 min after H<sub>2</sub>O<sub>2</sub> treatment, causing a 25-fold increase in the phospho-ERK-2/ERK-2 ratio, and gradually declined over 2 hr. One-way ANOVA for the H<sub>2</sub>O<sub>2</sub> time course yielded a calculated *F* value of 3.12 (*p* = 0.049; Figure 2). The 30-min time point was used for further experiments evaluating

the concentration-dependent activation of ERK-2. H<sub>2</sub>O<sub>2</sub>-induced ERK-2 activation was maximal at 10  $\mu$ M H<sub>2</sub>O<sub>2</sub> and did not significantly increase further at 50  $\mu$ M H<sub>2</sub>O<sub>2</sub> (Figure 3). One-way ANOVA for the H<sub>2</sub>O<sub>2</sub> dose response yielded a calculated *F* value of 3.85 (*p* = 0.038). In all of these experiments, the signal for total ERK-2 protein did not change with H<sub>2</sub>O<sub>2</sub> treatment.

**Delayed activation of ERK-2 by LPS.** We then investigated whether ERK-2 in XTC-2 cells was a target for endotoxin-mediated cell stress. ERK-2 phosphorylation peaked 60 min after LPS treatment, causing a 5- to 10-fold increase in the phospho-ERK-2/ERK-2 ratio (Figure 4). These data indicated that LPS was a weaker activator of ERK-2 compared to H<sub>2</sub>O<sub>2</sub>, and the ERK-2 phosphorylation response to LPS was delayed by as much as 30 min compared to H<sub>2</sub>O<sub>2</sub>. LPS caused a concentration-dependent increase in ERK-2 activation that was

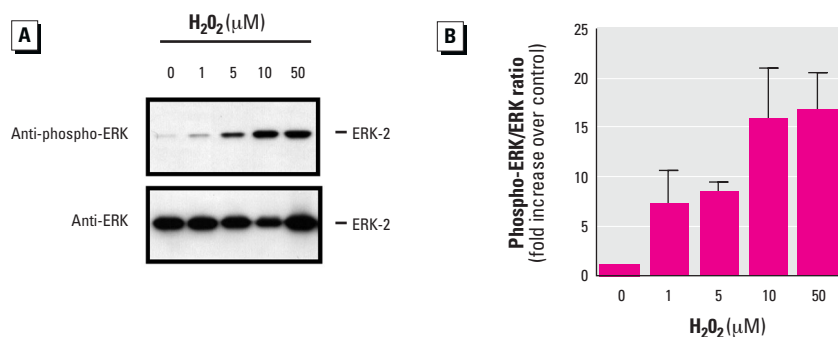
maximal at 50–500  $\mu$ g/mL LPS (Figure 5). One-way ANOVA yielded a calculated *F* value of 2.57 (*p* = 0.08) for the LPS time-course data (Figure 4) and of 2.80 (*p* = 0.08) for dose response (Figure 5).

**Effect of ERK inhibition on H<sub>2</sub>O<sub>2</sub>- and LPS-induced cytotoxicity.** H<sub>2</sub>O<sub>2</sub> and LPS both caused a concentration-dependent increase in cytotoxicity after 24 hr as determined by MTT assay (Figure 6). Parallel cultures of cells in log-phase growth (i.e., cells exposed in medium containing 10% fetal bovine serum) were pretreated with either 50  $\mu$ M PD98059 (MEK/ERK inhibitor) or dimethylsulfoxide vehicle control for 1 hr before treatment with H<sub>2</sub>O<sub>2</sub> or LPS. Pretreatment of XTC-2 cells with PD98059 significantly reduced the cytotoxic response to either H<sub>2</sub>O<sub>2</sub> or LPS (Figure 6). Marginal cytotoxicity was observed in quiescent cultures of XTC-2 cells treated with H<sub>2</sub>O<sub>2</sub> or LPS, indicating that co-stimulation of cells with growth-stimulatory factors in serum plus the environmental stressor (H<sub>2</sub>O<sub>2</sub> or LPS) were required for a maximal cytotoxic response. These data indicated that the cytotoxic effects of reactive oxygen species or bacterial endotoxin are mediated at least in part through activation of the ERK pathway. Significant differences between control versus treatment with PD98059 were determined by paired *t*-tests.

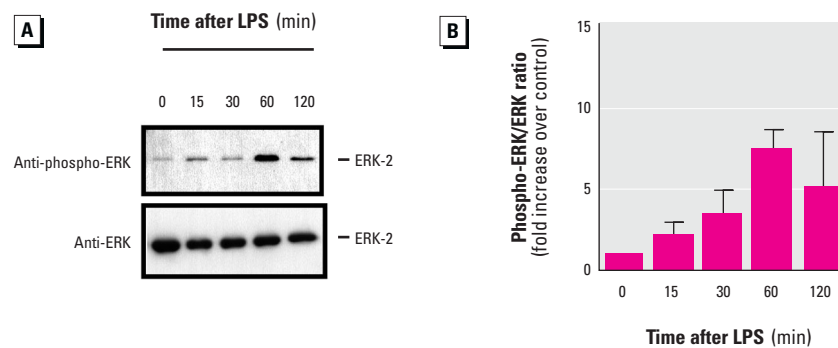
## Discussion

In this study, we found that the ERK-2 MAP kinase pathway in *Xenopus* (XTC-2) cells is activated by oxidative stress (H<sub>2</sub>O<sub>2</sub>) or bacterial lipopolysaccharide (LPS). Moreover, ERK activation mediated, at least in part, the cytotoxic response to either H<sub>2</sub>O<sub>2</sub> or LPS. This *in vitro* model of MAP kinase activation could help identify pollutants in aquatic ecosystems that contribute to the problem of amphibian malformations and population decline. Amphibians have evolved strategies to cope with harsh changes in natural conditions, including dessication of temporary water sources (24). Despite this resilience in the face of natural environmental changes, amphibians may be particularly susceptible to a wide range of environmental toxicants that are present in polluted waters. The XTC-2 cell culture model might be useful for detecting cellular stress induced by pollutants that generate reactive oxygen species (e.g., metals, pesticides) or gram-negative bacteria that release endotoxin.

The XTC-2 *in vitro* model for measuring ERK-2 activation could also be used to test the effects of UV radiation generated in the laboratory. UV radiation has been proposed as a major stress factor for some amphibian populations (4,16,19). Moreover, it is important to consider that UV radiation



**Figure 3.** Phosphorylation of ERK-2 in XTC-2 cells occurs in a concentration-dependent manner by the reactive oxygen species, H<sub>2</sub>O<sub>2</sub>. Confluent, quiescent cultures of XTC-2 cells were treated with an increasing concentration of H<sub>2</sub>O<sub>2</sub> for 15 min before collecting cell lysates for Western blot analysis of phospho-ERK and ERK. (A) Representative Western blots showing that phosphorylation of ERK-2 is increased by H<sub>2</sub>O<sub>2</sub> maximally at 10  $\mu$ M (upper frame), whereas the level of unphosphorylated ERK does not change (lower frame). (B) Relative levels of ERK-2 activation determined by densitometric scanning of the phospho-ERK bands and normalized to the ERK signal. Data are the means  $\pm$  SEM of three separate experiments and are presented as the phospho-ERK-2/ERK-2 ratio. One-way ANOVA yielded a calculated *F* value of 3.85 (*p* = 0.038).



**Figure 4.** Phosphorylation of ERK-2 in XTC-2 cells by the bacterial endotoxin LPS in a time-dependent manner. Confluent, quiescent XTC-2 cells were treated with 100  $\mu$ g/mL LPS for 15, 30, 60, and 120 min. (A) Representative Western blot experiment showing that maximum phosphorylation of ERK-2 was observed at 60 min after treatment with LPS. (B) Relative levels of ERK-2 activation determined by densitometric scanning of the phospho-ERK bands and normalized to the ERK signal. Data are the means  $\pm$  SEM of three separate experiments and presented as the phospho-ERK-2/ERK-2 ratio. One-way ANOVA yielded a calculated *F* value of 2.57 (*p* = 0.08).



may alter the chemical structure of some xenobiotics and thereby enhance toxicity. For example, Zaga and colleagues (25) reported that UV radiation enhanced the toxicity of carbaryl (a broad spectrum insecticide) to *Xenopus laevis* and *Hyla versicolor* embryos and tadpoles. Therefore, future investigation should include studies on agents such as carbaryl that are photoactivated by UV radiation to determine if this enhances the cellular stress response (e.g., ERK-2 activation) in cultured *Xenopus* cells.

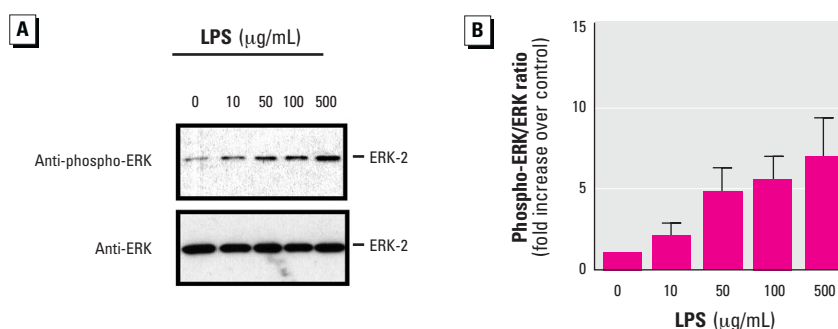
The activation of MAP kinase signaling pathways in response to endogenous growth factors is well established and produces cell growth or differentiation (26,27). A variety of polypeptide growth factors (e.g., fibroblast growth factor-2 (FGF), platelet-derived growth factor (PDGF), epidermal growth factor (EGF) bind to specific cell-surface receptors and initiate an intracellular signal transduction cascade involving the sequential phosphorylation of Ras, Raf, Map kinase kinase (MEK), and ERK. ERK then translocates to the nucleus, where it phosphorylates a variety of transcription factors that control gene expression (9). *Xenopus* oocyte maturation requires phosphorylation and activation of p42 MAP kinase, and the corresponding dephosphorylation and inactivation of the enzyme is crucial for the transition of the zygote from meiosis through the first phases of mitosis (10). Likewise, Bitangcol et al. (28) found that p42 MAP kinase activation regulates the normal progression through the cell cycle and is an essential component of the spindle assembly checkpoint. Furthermore, during *Xenopus* development, ordinary mesoderm differentiation relies on MAP kinase activation in response to FGF. However, overexpression of *Xenopus* MAP kinase phosphatase obstructs mesoderm induction by FGF and produces characteristic defects in mesoderm formation in intact embryos (29). We postulate that oxidative or bacterial stress that activates the ERK-2 pathway during development could contribute to the morphologic anomalies seen in frogs, especially those involving mesodermal derivatives such as bone and muscle differentiation in the limbs. In fact, our cytotoxicity data showed that H<sub>2</sub>O<sub>2</sub> or LPS caused more toxicity when cells were exposed in log-phase growth (i.e., in the presence of medium containing 10% fetal bovine serum) compared to cells in a quiescent state (Figure 6). Cells undergoing mitosis appear to be more susceptible to environmental stress.

Although the ERK-2 pathway is essential for growth and development mediated by growth factors, increasing evidence suggests that the ERK pathway is the target of environmental stress. Several environmental stressors (oxidants, metals, or UV radiation)

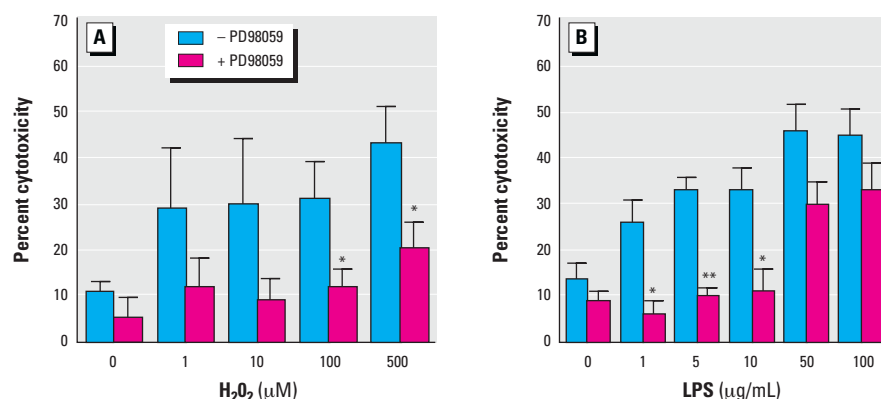
cause ERK activation by first causing phosphorylation of the intracellular domain of the EGF receptor in a ligand-independent manner (i.e., in the absence of EGF) (9,15,30). ERK activation by these environmental agents generally causes cell death (apoptosis) rather than normal cell growth and differentiation. For example, H<sub>2</sub>O<sub>2</sub> causes a cytotoxic response of mammalian connective tissue cells via the EGF receptor pathway leading to ERK activation, and a metabolic inhibitor of the ERK pathway (PD98059) blocked this cytotoxic response (23,30). We also observed that pretreatment of XTC-2 cells with PD98059 reduced the cytotoxic response of these cells to either H<sub>2</sub>O<sub>2</sub> or LPS (Figure 6). Therefore, these environmental agents appear to cause XTC-2 cell death in part via activation of ERK-2.

Our data show that both oxidative and bacterial stress trigger ERK phosphorylation in a time- and concentration-dependent manner. H<sub>2</sub>O<sub>2</sub> caused a rapid and more

pronounced phosphorylation of ERK-2, because maximal phosphorylation took place at 30 min (Figure 2) with a 20-fold increase from the control (Figure 3). In contrast, LPS was a weaker activator of ERK-2. The maximum phosphorylation response occurred after 60 min (Figure 4) with only a 5- to 10-fold increase from the control (Figure 5). These differences are likely due to different mechanisms of ERK activation by H<sub>2</sub>O<sub>2</sub> and LPS. As mentioned above, H<sub>2</sub>O<sub>2</sub> crosses the cell membrane and activates the intracellular domain of the EGF receptor, resulting in downstream activation of Ras, Raf, MEK, and ERK (30). In contrast, LPS docks to a cell-surface receptor, mCD14, which then complexes with a transmembrane signaling receptor, the toll-like receptor 4 (TLR4) (31). Activation of TLR4 then leads to ERK activation through a mechanism that is not yet clear (32). The different time-course response of ERK-2 to H<sub>2</sub>O<sub>2</sub> in comparison to LPS could allow for discrimination of



**Figure 5.** Phosphorylation of ERK-2 in XTC-2 cells occurs in a concentration-dependent manner after treatment with LPS. Confluent, quiescent cultures of XTC-2 cells were treated with an increasing concentration of LPS for 30 min before collecting cell lysates for Western blot analysis of phospho-ERK and ERK. (A) Representative Western blots showing that phosphorylation of ERK-2 is increased by LPS maximally at 50 to 500 µg/mL (upper frame), whereas the level of unphosphorylated ERK does not change (lower frame). (B) Relative levels of ERK-2 activation determined by densitometric scanning of the phospho-ERK bands and normalized to the ERK signal. Data shown are the mean ± SEM of three separate experiments and are presented as the phospho-ERK-2/ERK-2 ratio. One-way ANOVA yielded a calculated *F* value of 2.79 (*p* = 0.08).



**Figure 6.** The cytotoxic response of XTC-2 cells after H<sub>2</sub>O<sub>2</sub> or LPS exposure is reduced by pretreatment with PD98059. Confluent cultures of XTC-2 cells in log-phase growth were pretreated with 50 µM PD98059 delivered in DMSO or with DMSO (vehicle control) alone for 1 hr before treatment of cells with an increasing concentration of H<sub>2</sub>O<sub>2</sub> or LPS. After 24 hr, an MTT cytotoxicity assay was performed as described in "Materials and Methods." Data shown are the mean ± SEM of three separate experiments. Significant differences between control and PD98059-treated cells were determined by paired *t*-test (\**p* < 0.05; \*\**p* < 0.01).

oxidative and bacterial stress in samples obtained from polluted waters.

Other types of pollution in aquatic systems should be considered as stress activators of MAP kinases. For example, metals may also represent a source of oxidative stress in polluted, aquatic environments (33). We have observed that vanadium pentoxide, a transition metal released from the industrial burning of fuel oil, causes ERK-2 activation in XTC-2 cells (data not shown). Also, pesticides have been associated with amphibian population declines (13). Certain pesticides recently have been reported to activate MAP kinase signaling pathways in a mammalian prostate cancer cell line (34). Therefore, it would be important to learn whether ERK-2 is activated by pesticides in the XTC-2 cell culture model.

In summary, we have shown that representative oxidative and bacterial agents cause activation of the ERK-2 MAP kinase in cultured *Xenopus* XTC-2 cells.  $H_2O_2$  and LPS caused different temporal and dose-response patterns of ERK-2 activation. Blocking ERK-2 activation with the MEK inhibitor PD98059 partially protected XTC-2 cells against the cytotoxic effects of  $H_2O_2$  or LPS. We propose that ERK-2 activation is a valuable biochemical marker for assessing factors that cause cellular stress in amphibians. Moreover, the XTC-2 cell culture model could be useful for rapidly assessing the "amphibian stress potential" of polluted waters. Future studies will address the utility of MAP kinase activation in XTC-2 cells as a screening assay for stressors present in the environment.

## REFERENCES AND NOTES

- Manuel JS. Malformed frogs: making the leap to humans. *Environ Health Perspect* 108:A31 (2000).
- Kaiser J. Deformed frogs leap into spotlight at health workshop. *Science* 278:2051–2052 (1997).
- Johnson PTJ, Lunde KB, Ritchie EG, Launer AE. The effect of trematode infection on amphibian limb development and survivorship. *Science* 284:802–804 (1999).
- Blaustein AR, Hoffman PD, Hokot DG, Kiesecker JM, Walls SC, Hays JB. UV repair and resistance to solar UV-B in amphibian eggs: a link to population declines? *Proc Natl Acad Sci USA* 91:1791–1795 (1994).
- Johnson NA. The case of the deformed frogs. *American Scientist* 88:362–363 (2000).
- Burkhardt JG, Ankley G, Bell H, Carpenter H, Fort D, Gardiner D, Gardner H, Hale R, Helgen JC, Jepson P et al. Strategies for assessing the implication of malformed frogs for environmental health. *Environ Health Perspect* 108:83–90 (2000).
- Blaustein AR, Wake, DB. Declining amphibian populations: a global phenomenon? *Trends Ecol Evol* 7:203–204 (1990).
- Pechmann JHK, Scott DE, Semlitsch RD, Caldwell JP, Vitt LJ, Gibbons WJ. Declining amphibian populations: the problem of separating human impacts from natural fluctuations. *Science* 253: 892–895 (1991).
- Schaeffer HJ, Weber MJ. Mitogen-activated protein kinases: specific messages from ubiquitous messengers. *Mol Cell Biol* 19:2435–2444 (1999).
- Sohaskey ML, Ferrell JE Jr. Distinct, constitutively active MAPK phosphatases function in *Xenopus* oocytes: implications for p42 MAPK regulation *in vivo*. *Mol Biol Cell* 10:3729–3743 (1999).
- Bonner JC, Rice AB, Lindroos PM, O'Brien PO, Dreher KL, Rosas I, Alfaro-Moreno E, Osornio-Vargas AR. Induction of the lung myofibroblast PDGF receptor system by urban ambient particles from Mexico City. *Am J Respir Cell Mol Biol* 19:672–680 (1998).
- Samet JM, Graves LM, Quay J, Dailey LA, Devlin RB, Ghio AJ, Wu W, Bromberg PA, Reed W. Activation of MAPKs in human bronchial epithelial cells exposed to metals. *Am J Physiol* 275:L551–L558 (1998).
- Sparling DW, Fellers GM, McConnell LL. Pesticides and amphibian population declines in California, USA. *Environ Toxicol Chem* 20:1591–1595 (2001).
- Yanase, H, Ando H, Horikawa M, Watanabe M, Mori T, Matsuda N. Possible involvement of ERK 1/2 in UVA-induced melanogenesis in cultured normal human epidermal melanocytes. *Pigment Cell Res* 14:103–109 (2001).
- Wang Y-Z, Bonner JC. Mechanism of extracellular signal-regulated kinase (ERK)-1 and ERK-2 activation by vanadium pentoxide in rat pulmonary myofibroblasts. *Am J Respir Cell Mol Biol* 22:590–596 (2000).
- Blaustein AR, Kiesecker JM, Chivers DP, Hokit DG, Marco AL, Belden LK, Hatch A. Effects of ultraviolet radiation on amphibians: field experiments. *Amer Zoologist* 38:799–812 (1998).
- Ankley GT, Tietge JE, Defoe DL, Jensen KM, Holcombe GW, Durham EJ, Diamond SA. Effects of ultraviolet light and methoprene on survival and development of *Rana pipiens*. *Environ Toxicol Chem* 17:2530–2542 (1998).
- Starnes SM, Kennedy CA, Petranka JW. Sensitivity of embryos of southern Appalachian amphibians to ambient solar UV-B radiation. *Conserv Biol* 14:277–282 (2000).
- Bruggeman DJ, Bantle JA, Goad C. Linking teratogenesis, growth, DNA photodamage to artificial ultraviolet B radiation in *Xenopus laevis* larvae. *Environ Toxicol Chem* 17:2114–2121 (1998).
- Fiskdal L, Tryland I, Nelis H. Rapid detection of coliform bacteria and influence of non-target bacteria. *Wat Sci Tech* 35:415–418 (1997).
- Hader D-P, Kumar HD, Smith RC, Worrest RC. Effects on aquatic ecosystems. *J Photochem Photobiol* 46:53–68 (1998).
- Yi S, Sullivan SK, Pitterle DM, Kennington EA, Graff JM, Blackshear PJ. Mechanisms of MARCKS gene activation during *Xenopus* development. *J Biol Chem* 272:29290–29300 (1997).
- Zhang P, Wang Y-Z, Kagan E, Bonner JC. Peroxynitrite targets the epidermal growth factor receptor, Raf-1, and MEK independently to activate MAPK. *J Biol Chem* 275:22479–22486 (2000).
- Bonner LA, Diehl WL, Altig RA. Physical, chemical and biological dynamics of five temporary dystrophic pools in central Mississippi. *Hydrobiologia* 353:77–89 (1997).
- Zaga A, Little EE, Rabeni CF, Ellersieck MR. Photoenhanced toxicity of a carbamate insecticide to early life stages of anuran amphibians. *Environ Toxicol Chem* 17:2543–2553 (1998).
- Cobb MH, Goldsmith EJ. How MAP kinases are regulated. *J Biol Chem* 270:14843–14846 (1995).
- Waskiewicz AJ, Cooper JA. Mitogen and stress response pathways: MAP kinase cascades and phosphatase regulation in mammals and yeast. *Curr Opin Cell Biol* 7:798–805 (1995).
- Bitangcol JC, Chau AS, Stadnick E, Lohka MJ, Dicken B, Shibuya EK. Activation of the p42 mitogen-activated protein kinase pathway inhibits Cdc2 activation and entry into M-phase in cycling *Xenopus* egg extracts. *Mol Biol Cell* 9:451–467 (1998).
- Umbhauer M, Marshall CJ, Mason CS, Old RW, Smith JC. Mesoderm induction in *Xenopus* caused by activation of MAP kinase. *Nature* 376:58–62 (1995).
- Guyton KZ, Liu Y, Gorospe M, Xu Q, Holbrook NJ. Activation of mitogen-activated protein kinase by  $H_2O_2$ : role in cell survival following oxidant injury. *J Biol Chem* 271:4138–4142 (1996).
- Means TK, Golenbock DT, Fenton MJ. The biology of toll-like receptors. *Cytokine Growth Factor Rev* 11:219–232 (2000).
- Yang H, Young DW, Gusovsky F, Chow JC. Cellular events mediated by lipopolysaccharide-stimulated toll-like receptor 4. *J Biol Chem* 275:20861–20866 (2000).
- Mielke HW, Gonzales CR, Smith MK, Mielke PW. Quantities and associations of lead, zinc, cadmium, manganese, chromium, nickel, vanadium, and copper in fresh Mississippi delta alluvium and New Orleans alluvial soils. *Sci Total Environ* 246:249–235 (2000).
- Tessier DM, Matsumura F. Increased ErbB2 tyrosine kinase activity, MAPK phosphorylation, and cell proliferation in the prostate cancer cell line LNCaP following treatment with select pesticides. *Toxicol Sci* 60:38–43 (2001).